

NOV 26 2003

SECTION 7

510(k) SUMMARY OF SAFETY EFFECTIVENESS

This 510(k) summary of safety and effectiveness information is being submitted in accordance with the requirements of the Safe Medical Device Act (SMDA) of 1990. The contents of this 510(k) summary have been provided in conformance with 21 CFR §807.92.

Date: October 22, 2003

Common/Usual Name: Pre-mixed Dialysate

Trade/Proprietary Name: NxStage PureFlow Solution

Classification Name: Hemodialysis systems and accessories (21 CFR 876.5820)

Device Classification: Class II

Product Code: 78 KPO – Dialysate Concentrate for Hemodialysis (Liquid or Powder)

Device Panel: Gastroenterology-Urology (GU)/Gastro-Renal (GRDB)

510(k) Sponsor & Owner/Operator: NxStage Medical, Inc
439 South Union St, Suite 501
Lawrence, MA 01843
Owner/Operator No. 9045797

Contact Person: Norma LeMay
Regulatory Affairs Manager

Device Description:
The NxStage PureFlow Solutions are sterile, non-pyrogenic dialysis solutions provided in single use flexible PVC bags varying in sizes from 1000 to 5000ml. The dialysis solutions are intended for use with renal replacement therapy systems that utilize sterile premixed dialysate.

Substantial Equivalence:

This submission is a Special 510(k) Device Modification as described in FDA's guidance document entitled "The New 510(k) Paradigm - Alternate Approaches to Demonstrating Substantial Equivalence in Pre-Market Notifications." In support of this 510(k), NxStage Medical has provided certification of compliance to 21 CFR 820.30 Design Control requirements. Design validation testing has been performed to ensure that the modified device meets design specifications. The modified NxStage PureFlow Solution has been compared to the baseline as cleared in K022913 and found to be substantial equivalent.

Conclusion:

Based on the device indications for use, comparison of descriptive and technological characteristics, and design control certification, the modified NxStage PureFlow Solution has been shown to meet the minimum requirements that are considered acceptable for its intended use.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 26 2003

Ms. Norma LeMay
Regulatory Affairs Manager
NxStage Medical, Inc.
439 South Union Street, Suite 501
LAWRENCE MA 01843

Re: K033386

Trade/Device Name: NxStage PureFlow Solution
Regulation Number: 21 CFR §876.5820
Regulation Name: Hemodialysis system and accessories
Regulatory Class: II
Product Code: 78 KPO
Dated: October 22, 2003
Received: October 27, 2003

Dear Ms. LeMay:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of the letter:

8xx.1xxx	(301) 594-4591
876.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4616
884.2xxx, 3xxx, 4xxx, 5xxx, 6xxx	(301) 594-4616
892.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4654
Other	(301) 594-4692

Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97) you may obtain. Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>.

Sincerely yours,



Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

INDICATIONS FOR USE STATEMENT

Page 1 of 1

510(k) Number (if known): K033386

Device Name: NxStage PureFlow Solution

Indications for Use:

The NxStage Pureflow Solution is indicated for use with renal replacement therapy systems that utilize sterile premixed dialysate.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓
(Per 21 CFR 801.109)

OR

Over-the-Counter Use

David G. Flynn
(Division Sign-Off)

Division of Reproductive, Abdominal,
and Radiological Devices

510(k) Number K033386

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